Centers for Medicare & Medicaid Services

[Document Identifiers: CMS-10137, CMS-10141, 10773 and 10494]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish notice in the *Federal Register* concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments on the collection(s) of information must be received by the OMB desk officer by [INSERT DATE 30 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER].

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain . Find this particular information collection by selecting "Currently under 30-day Review - Open for Public Comments" or by using the search function.

To obtain copies of a supporting statement and any related forms for the proposed

collection(s) summarized in this notice, you may make your request using one of following:

 Access CMS' Web Site address at Web Site address at: https://www.cms.gov/Regulations-and-

Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html

FOR FURTHER INFORMATION CONTACT: William Parham at (410) 786-4669.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501-3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term "collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-day notice in the *Federal Register* concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. Type of Information Collection Request: Revision of a currently approved collection; Title of Information Collection: Solicitation for Applications for Medicare Prescription Drug Plan 2023 Contracts; Use: Coverage for the prescription drug benefit is provided through contracted prescription drug plans (PDPs) or through Medicare Advantage (MA) plans that offer integrated prescription drug and health care coverage (MA-PD plans). Cost Plans that are regulated under Section 1876 of the Social Security Act, and Employer Group Waiver Plans (EGWP) may also provide a Part D benefit. Organizations wishing to provide services under the Prescription Drug Benefit Program must complete an application, negotiate rates, and receive final approval from CMS. Existing Part D Sponsors may also expand their contracted service area by completing the Service Area Expansion (SAE) application.

Collection of this information is mandated in Part D of the Medicare Prescription Drug, Improvement, and Modernization Act of 2003 (MMA) in Subpart 3. The application requirements are codified in Subpart K of 42 CFR 423 entitled "Application Procedures and Contracts with PDP Sponsors."

The information will be collected under the solicitation of proposals from PDP, MA-PD, Cost Plan, Program of All Inclusive Care for the Elderly (PACE), and EGWP applicants. The collected information will be used by CMS to: (1) ensure that applicants meet CMS requirements for offering Part D plans (including network adequacy, contracting requirements, and compliance program requirements, as described in the application), (2) support the determination of contract awards. *Form Number:* CMS-10137 (OMB control number: 0938-0936); *Frequency: Yearly; Affected Public:* Businesses or other for-profits, Not-for-profit institutions; *Number of Respondents:* 716; *Total Annual Responses:* 382; *Total Annual Hours:* 1,716. (For policy questions regarding this collection contact Arianne Spaccarelli at 410-786-5715.)

- 2. Type of Information Collection Request: Revision of a currently approved collection; Title of Information Collection: Medicare Prescription Drug Benefit Program; Use: Plan sponsor and State information is used by CMS to approve contract applications, monitor compliance with contract requirements, make proper payment to plans, and ensure that correct information is disclosed to potential and current enrollees. Form Number: CMS-10141 (OMB control number: 0938-0964); Frequency: Once; Affected Public: Private sector (Business or other for-profit and Not-for-profit institutions); Number of Respondents: 11,771,497; Total Annual Responses: 675,231,213; Total Annual Hours: 9,312,314. (For policy questions regarding this collection contact Maureen Connors at 410-786-4132.)
- 3. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Non-Quantitative Treatment Limitation Analyses and Compliance Under MHPAEA; *Use:* The Paul Wellstone and Pete Domenici Mental Health Parity and Addiction Equity Act of 2008 (MHPAEA) (P.L.110-343) generally requires that

group health plans and group health insurance issuers offering mental health or substance use disorder (MH/SUD) benefits in addition to medical and surgical (med/surg) benefits do not apply any more restrictive financial requirements (e.g., co-pays, deductibles) and/or treatment limitations (e.g., visit limits, prior authorizations) to MH/SUD benefits than those requirements and/or limitations applied to substantially all med/surg benefits. The Patient Protection and Affordable Care Act, Pub. L. 111-148, was enacted on March 23, 2010, and the Health Care and Education Reconciliation Act of 2010, Pub. L. 111-152, was enacted on March 30, 2010. These statutes are collectively known as the "Affordable Care Act." The Affordable Care Act extended MHPAEA to apply to the individual health insurance market. MHPAEA does not apply directly to small group health plans, although its requirements are applied indirectly in connection with the Affordable Care Act's essential health benefit requirements. The Consolidated Appropriations Act, 2021 (the Appropriations Act) was enacted on December 27, 2020. The Appropriations Act amended MHPAEA, in part, by expressly requiring group health plans and health insurance issuers offering group or individual health insurance coverage that offer both med/surg benefits and MH/SUD benefits and that impose non-quantitative treatment limitations (NQTLs) on MH/SUD benefits to perform and document their comparative analyses of the design and application of NQTLs. Further, beginning 45 days after the date of enactment of the Appropriations Act, group health plans and health insurance issuers offering group or individual health insurance coverage must make their comparative analyses available to the Departments of Labor, Health and Human Services (HHS), and the Treasury or applicable state authorities, upon request. The Secretary of HHS is required to request the comparative analyses for plans that involve potential violations of MHPAEA or complaints regarding noncompliance with MHPAEA that concern NQTLs and any other instances in which the Secretary determines appropriate. The Appropriations Act also requires the Secretary of HHS to submit to Congress, and make publicly available, an annual report on the conclusions of the reviews. Form Number: CMS-10773 (OMB control number: 0938-1393); Frequency: On Occasion; Affected Public:

State, Local, or Tribal Governments, Private Sector; Number of Respondents: 250,137; Total

Annual Responses: 36,461; Total Annual Hours: 1,013,184. (For policy questions regarding this

collection, contact Usree Bandyopadhyay at 410-786-6650.)

Type of Information Collection Request: Revision of a currently approved 4.

collection; Title of Information Collection: Exchange Functions: Standards for Navigators and

Non-Navigator Assistance Personnel-CAC; *Use*: Section 1321(a)(1) of the Affordable Care Act

directs and authorizes the Secretary to issue regulations setting standards for meeting the

requirements under title I of the Affordable Care Act, with respect to, among other things, the

establishment and operation of Exchanges. Pursuant to this authority, regulations establishing the

certified application counselor program have been finalized at 45 CFR 155.225. In accordance

with 155.225(d)(1) and (7), certified application counselors in all Exchanges are required to be

initially certified and recertified on at least an annual basis and successfully complete Exchange

required training. Form Number: CMS-10494 (OMB control number: 0938-1205); Frequency:

On Occasion; Affected Public: State, Local, or Tribal Governments, Private Sector (not-for-profit

institutions); individuals or households; Number of Respondents: 278,072; Total Annual

Responses: 278,072; Total Annual Hours: 918,024. (For policy questions regarding this

collection contact Evonne Muoneke at 301-492-4402.)

Dated: October 21, 2021.

William N. Parham, III,

Director,

Paperwork Reduction Staff,

Office of Strategic Operations and Regulatory Affairs.

4120-01-U-P

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